

Notice of References Cited	Application/Control No. 10/021,955	Applicant(s)/Patent Under Reexamination LUPSKI ET AL.	
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U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
	A	US-			
	B	US-			
	C	US-			
	D	US-			
	E	US-			
	F	US-			
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FOREIGN PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Country	Name	Classification
	N					
	O					
	P					
	Q					
	R					
	S					
	T					

NON-PATENT DOCUMENTS

*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
*	U	Roa et al. Dejerine-Scottas syndrome associated with point mutation in the peripheral myelin protein 22 (PMP22) gene. Nature Genetics, Vol. 5, pp. 269-273, 1993.
*	V	Guilbot et al. A mutation in periaxin is responsible for CMT4F, an autosomal recessive form of Charcot-Marie-Tooth disease. Human Mole.Genet., Vo. 10, No. 4, pp. 415-421, 2001.
	W	
	X	

*A copy of this reference is not being furnished with this Office action. (See MPEP § 707.05(a).)
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U.S. PATENT DOCUMENTS

*		Document Number Country Code-Number-Kind Code	Date MM-YYYY	Name	Classification
*	A	US- 5,691,144	11-1997	Boss et al-	435/6
*	B	US- 5,780,223	7-1998	Lupski et al-	435/6
	C	US-			
	D	US-			
	E	US-			
	F	US-			
	G	US-			
	H	US-			
	I	US-			
	J	US-			
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	O					
	P					
	Q					
	R					
	S					
	T					

NON-PATENT DOCUMENTS

*		Include as applicable: Author, Title Date, Publisher, Edition or Volume, Pertinent Pages)
	U	Gillespie et al. The gene encoding the schwann cell protein periaxin localizes on mouse chromosome 7 (PRX). Geneomics, Vo 41, pages 297-298, 1997.
	V	Boerkoel et al. Periaxin mutations cause recessive Dejerine-Sottas Neuropathy. Am. J. Hum. Genet., Vol. 68, pages 325-333, 2001.
*	W	Timmerman et al. Novel missense mutation in the early growth response 2 gene associated with Dejerine-Sttas syndrome phenotype. Neurology, Vol. 52, pp. 1827-1832, 1999.
*	X	Gillespie et al. Periaxin, a novel protein of myelinating schwann cells with a possible role in axonal ensheathment. Neuron, Vol 12, pp. 497-508, 1994.

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Dates in MM-YYYY format are publication dates. Classifications may be US or foreign.